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AMENDMENTS TO THE CLAIMS:

Claim 1. (Previously Presented) A pharmaceutical composition consisting essentially of fexofenadine or a pharmaceutical acceptable acid addition salt thereof, about 10 wt. % to about 70 wt. % of lactose, and about 1 wt. % to about 40 wt. % of a low-substituted hydroxypropyl cellulose, wherein the weight percents are based on the total weight of the pharmaceutical composition.

Claim 2. (Original) The composition according to Claim 1, wherein the salt of fexofenadine is fexofenadine hydrochloride.

Claim 3. (Original) The composition according to Claim 1, wherein the amount of fexofenadine or pharmaceutical acceptable salt thereof is from about 1 wt. % to about 80 wt. %, based on the total weight of the pharmaceutical composition.

Claim 4. (Original) The composition according to Claim 3, wherein the amount of fexofenadine or pharmaceutical acceptable salt thereof is from about 5 wt. % to about 50 wt. %, based on the total weight of the pharmaceutical composition.

Claim 5. (Original) The composition according to Claim 4, wherein the amount of fexofenadine or pharmaceutical acceptable salt thereof is from about 20 wt. % to about 35 wt. %, based on the total weight of the pharmaceutical composition.

Claim 6. (Original) The composition according to Claim 1, wherein the amount of fexofenadine or pharmaceutical acceptable salt thereof is from about 10 mg to about 200 mg.

Claim 7. (Original) The composition according to Claim 6, wherein the amount of fexofenadine or pharmaceutical acceptable salt thereof is from about 30 mg to about 180 mg.

Claim 8. (Original) The composition according to Claim 1, wherein the lactose is selected from the group consisting of lactose monohydrate, lactose anhydrous, α-lactose, ß-lactose, and combinations thereof.

Claim 9. (Original) The composition according to Claim 8, wherein the lactose is lactose monohydrate.